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(54) **Feed for domestic fowls.**

(57) A feed for domestic fowls which comprises an indigestible dextrin prepared by roasting a starch in the presence of a mineral acid to give a pyrodextrin and treating the pyrodextrin with α -amylase or an indigestible dextrin prepared by further treating the α -amylase-treated product with glucoamylase, followed by filtration, purification and separation and removal of digestible fractions of the product through chromatography using an ion exchange resin. The feed serves to inhibit the formation of fatty liver, to improve the meat quality and the rate of egg laying, to improve the eggshell strength and to reduce the cholesterol content in the egg.

BACKGROUND OF THE INVENTION

The present invention relates to a feed for domestic fowls and more specifically to a feed for domestic fowls which can inhibit the formation of fatty liver and can in turn improve the meat quality and which permits the reduction in the content of cholesterol in eggs and can improve the egg-laying rate and the eggshell strength.

It has been a recent tendency to make the formula feeds for fowls more thick (so-called thickened feeds) or increase the content of proteins and make them calorific for the improvement of the efficiency thereof. Thus, fowls take energy more than the required energy intake and accordingly, there is a sign of frequent incidence of abnormalities in lipometabolism such as excess accumulation of fats in, for instance, peritoneal cavity, intestinal tracts and hypodermis or formation of fatty liver. This accordingly leads to reduction in the performance of meat production (inclusive of viscus meat) in case of the chicken breed and reduction in the egg-laying rate and in the rate of cracked eggs due to reduction in the eggshell strength in case of the egg breed.

Recently, eating habits have been improved and on the other hand, the diseases of adult people have become a matter of great concern, in particular, arterial scleroses have widely and deeply been recognized. Cholesterols derived from foods have been known as causative substances of arterial scleroses. Recent investigations have proved that there is a positive correlation between the quantity of cholesterols taken in the form of foods and the content thereof in the serum. Moreover, it has also been recognized that a positive correlation is observed between the quantity of cholesterols taken in the form of foods and the mortality rate of cardiopathy due to arterial scleroses and it has been proved that the mortality rate of cardiopathy due to arterial scleroses increases as the amount of cholesterols taken in the form of foods increases.

The fowl egg is a typical example of foods having high contents of cholesterols. The amounts of various foods taken and the mortality rate of cardiopathy were examined on adult male persons of 55 to 59-year-old and it was proved that there was a significant positive correlation between the amount of fowl egg-intake and the mortality rate of cardiopathy ($r = 0.666$). Nevertheless, the fowl egg is an excellent nutrient substance as a source of high quality protein, is a food desirably taken at least one per day and accordingly, there has been desired for the development of a technique capable of reducing the content of cholesterols in the fowl egg.

Under such circumstances, there has been proposed an attempt which comprises adding, for instance, middle chain fatty acids, unsaturated fatty acids, amino acids and/or vitamins to feeds for domestic fowls in order to prevent accumulation of fats in the fowl body and to thus inhibit the formation of fatty liver. However, most of these substances are expensive and are not practically used as additives for the feed. Furthermore, there is a strong suspicion that chemical substances remaining in the meat and eggs of fowls may adversely affect the human body. Accordingly, the use of, in particular, synthetic chemical substances is not preferred from the viewpoint of such problem of safety. Thus, there has been desired for the development of cheap and effective feeds for domestic fowls.

SUMMARY OF THE INVENTION

Accordingly, an object of the present invention is to provide a feed for domestic fowls which is effective over the entire breeding period of the fowl, and which can prevent, in particular, accumulation of fats in the fowl body and the formation of fatty liver to thus improve the meat quality, to reduce the content of cholesterols in eggs and to improve the egg-laying rate and the eggshell strength.

The inventors of this invention have conducted various studies to solve the foregoing problems associated with the conventional feeds for domestic fowls, have found out that the use of indigestible dextrin is quite effective for the improvement of the feeds and thus have completed the present invention.

Accordingly, the present invention relates to a feed for domestic fowls which comprises an indigestible dextrin, wherein the indigestible dextrin may be those prepared by roasting a starch in the presence of a mineral acid to give a pyrodextrin and treating the pyrodextrin with α -amylase; those prepared by treating the foregoing indigestible dextrin, treated with α -amylase, with glucoamylase, filtering and purifying the product, and then separating and removing a digestible fraction from the product through chromatography with an ion exchange resin; and those commercially available as will be detailed below.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

The feed for domestic fowls comprising an indigestible dextrin of the present invention will be detailed below.

In the preparation of the indigestible dextrin used in the invention, a starch is first heat-treated in the presence of a mineral acid to give a pyrodextrin. Starches as raw materials for the indigestible dextrin incorporated into the feed for domestic fowls according to the present invention are not restricted to particular ones, but

specific examples thereof include those derived from corn, potato, sweet potato, tapioca, wheat, barley and rice.

More specifically, a starch is mixed with a mineral acid (such as hydrochloric acid, nitric acid or sulfuric acid), preferably hydrochloric acid in an amount, for instance, ranging from 3 to 10 parts by weight of a 1% by weight aqueous hydrochloric acid solution per 100 parts by weight of the starch used and the resulting mixture is heated to give a pyrodextrin as an intermediate product. Preferably, the aqueous solution of the starch and the mineral acid is uniformly mixed, prior to the heat treatment, by stirring and aging in an appropriate mixer and then pre-drying at a temperature on the order of 100 to 120 °C to reduce the moisture content of the mixture to approximately 5% by weight. The heat treatment is carried out at a temperature ranging from 150 to 220 °C for 10 minutes to 5 hours, preferably 15 to 60 minutes unlike the heating conditions for the heat treatment of the conventional dextrans (white dextrin and yellow dextrin) in the presence of acids. In this respect, the higher the temperature during the heat-treatment, the higher the content of indigestible components in the resulting product. However, the amount of colored substances in the product starts to increase at a temperature of about 180 °C. Thus, the heat treatment is more preferably performed at a temperature ranging from 150 to 180 °C.

Alternatively, it is also possible to carry out the heat treatment at an elevated temperature for a short period of time by properly selecting a heating apparatus to be used. For instance, the mixture can efficiently and uniformly be heat-treated in an apparatus such as an extruder. It is desirable to properly control the conditions for the heat-treatment while taking into consideration the amount of the starch powder to be heat-treated and the content of the indigestible components required for a particular product.

Then the pyrodextrin thus obtained is dissolved in water to give an aqueous solution having a concentration ranging from 30 to 50% by weight. The solution is hydrolyzed by treating with α -amylase. Any known α -amylase product can be used, but the most preferred example of commercially available α -amylase is Termamyl® (available from Novo Nordisk Bioindustry Company; heat-resistant α -amylase produced by a strain *Bacillus licheniformis*).

The aqueous solution of the pyrodextrin is acidic in nature due to the mineral acid added during roasting and therefore, the pH of the aqueous solution of the pyrodextrin is controlled to the optimum pH of α -amylase. Any basic compound commonly known can be used for controlling the pH, but a sodium hydroxide solution is most effective. The pH value preferably ranges from 5.5 to 6.5. If it is less than the lower limit, the reaction rate is reduced, while if it exceeds the upper limit, the product is markedly colored. Preferably, α -amylase is added to the solution after the pH adjustment and it is added thereto in an amount ranging from 0.05 to 0.2% by weight on the basis of the amount of the pyrodextrin.

The hydrolysis with α -amylase proceeds at a temperature of about 80 to 100 °C, but preferably 85 to 90 °C, since coloring of the product becomes significant at a high temperature. The reaction time in general ranges from 5 minutes to 3 hours, for instance, a satisfactory result can be ensured by carrying out the reaction for about one hour.

The hydrolyzed solution thus obtained generally contains indigestible components in an amount ranging from 40 to 60% by weight on the basis of the weight of the solid contents of the solution. The term "indigestible dextrin" herein used means the dextrin comprising, as principal components, indigestible components and includes, for instance, a liquid product obtained by concentrating the foregoing hydrolyzed solution per se, a powdery product obtained by drying the solution through, for instance, spray drying, those obtained by optionally subjecting the hydrolyzed solution to purification treatments such as decoloring with, for instance, active carbon, filtration and desalting and then concentrating into a liquid product or drying to give a powdery product, or those obtained by further subjecting these products to a treatment with glucoamylase as will be detailed below to increase the content of the indigestible components. In this respect, the product obtained through only the α -amylase treatment and having a content of the indigestible components ranging from 40 to 60% by weight on the basis of the weight of the solid contents thereof will hereunder be referred to as "indigestible dextrin A".

As has been discussed above, the foregoing indigestible dextrin, i.e., the indigestible dextrin A can be additionally hydrolyzed with glucoamylase to further increase the content of the indigestible components. The glucoamylase is not restricted to those derived from specific origins and thus any commercially available ones may effectively be used in the invention. Moreover, glucoamylase products commonly used in general comprise a small amount of α -amylase. Therefore, the pyrodextrin can be directly treated with glucoamylase comprising a small amount of α -amylase without treating with α -amylase to give the indigestible dextrin as the effective component of the present invention. However, if the glucoamylase used contains only a trace amount of α -amylase, the resulting product has only a low content of the indigestible components. Therefore, it is most preferred to treat the pyrodextrin with α -amylase then with glucoamylase. The glucoamylase treatment is preferably carried out at a pH ranging from 5.0 to 6.0. The amount of glucoamylase used in the treatment suitably ranges from 0.05 to 0.2% by weight on the basis of the weight of the indigestible dextrin as in the case of the

α -amylase treatment. The hydrolysis is performed at a temperature ranging from about 55 to 60°C and it is in general sufficient to carry out the hydrolysis for 24 to 48 hours.

Incidentally, it is evident to those skilled in the art that the amounts of α -amylase and glucoamylase are not restricted to the ranges defined above and that the amounts thereof can arbitrarily be adjusted depending on the titers of the amylase products used. Moreover, the reaction time can arbitrarily be adjusted through the control of the amounts thereof to be added. In addition, the filtration rate during the purification can be improved by hydrolyzing the pyrodextrin with α -amylase, then steaming the hydrolyzed solution under pressure at a temperature ranging from 115 to 135 °C, again treating with α -amylase and finally treating with glucoamylase.

After treating with glucoamylase, the glucoamylase is deactivated by reducing the pH of the solution to about 3.5 or elevating the temperature of the solution up to about 80 °C. Thereafter, the solution is optionally subjected to decoloring with the usual active carbon, filtration, and desalting and decoloring with an ion exchange resin. Then the solution is concentrated to a concentration on the order of 50% by weight, followed by chromatography using an ion exchange resin and separation and removal of the resulting glucose. In this case, strongly acidic cation exchange resins can widely be used.

Specific examples of preferred strong acidic cation exchange resins include Amberlite IR-116, IR-118, IR120-B, XT-1022E and XT-471F (trade names of the products available from Organo Co., Ltd.), Diaion 2K-18, SKK-102, SK-104, SK-106, SK-110, SK-112, SK-116 and FR-01 (trade names of the products available from Mitsubishi Chemical Industries Ltd.), XFS-43281.00, 43280.00, 43279.00 and 43278.00 (trade names of the products available from Dow Chemical Japan Ltd.).

In general, these resins are preferably converted into alkali metal or alkaline earth metal-forms prior to practical use. To improve the separability of the indigestible components from the glucose component, the flow rate of the solution passing through a column is preferably adjusted depending on the kinds of resins used, but the flow rate (SV) in general ranges from 0.1 to 0.6, preferably 0.2 to 0.4. The use of a flow rate beyond the range defined above may impair the workability and separability. The temperature during passing the solution through the column ranges from 20 to 70 °C and preferably 50 to 70°C. The use of a temperature of less than the lower limit may impair the separability of the indigestible component-containing fraction from the glucose-containing fraction or the viscosity of the solution increases and the capacity of the resin is in turn impaired. On the other hand, if the temperature exceeds the upper limit, the solution causes browning and other properties thereof are often impaired.

The indigestible dextrin obtained by additionally treating the indigestible dextrin A with glucoamylase and having an indigestible component-content of, in general, 50 to 90% by weight on the basis of the weight of the solid contents will hereunder be referred to as "indigestible dextrin B".

The indigestible dextrin used in the invention as the effective component may be commercially available ones. Examples of such easily and commercially available indigestible dextrin products include "Pinefiber" (trade name of the product available from Matsutani Chemical Industries Co., Ltd.) for the foregoing indigestible dextrin A obtained by treating the pyrodextrin simply with α -amylase and "Fibersol #2 (trade name of the product available from Matsutani Chemical Industries Co., Ltd.) for the foregoing indigestible dextrin B obtained by treating the pyrodextrin with α -amylase and then with glucoamylase. These commercially available indigestible dextrins comprise the indigestible components in amounts of about 55% by weight and about 90% by weight respectively and both of them have an average molecular weight of about 1600. Conventional formula of feeds for domestic fowls comprise crude fibers in the form of indigestible polysaccharides in an amount ranging from about 3 to 6% by weight, but the crude fibers are insoluble in water. On the other hand, the indigestible dextrin used in the invention as the effective component of the feed for domestic fowls is soluble in water and substantially differs from the conventionally known indigestible polysaccharides.

The amount of the indigestible dextrin is preferably not less than 0.2% by weight for the indigestible dextrin A and not less than 0.1% by weight for the indigestible dextrin B, on the basis of the total weight of the feed for domestic fowls. If the amount of the indigestible dextrin A or the indigestible dextrin B is less than 0.2% by weight or less than 0.1% by weight respectively, the resulting feed does not satisfactorily exhibit the desired effects of the present invention. Moreover, if the amount of the indigestible dextrin A or the indigestible dextrin B is more than 10% by weight or more than 5% by weight respectively, any further enhancement of the effects through the addition thereof is not expected. Therefore, the amount of the indigestible dextrin A preferably ranges from 0.2 to 10% by weight and that of the indigestible dextrin B preferably ranges from 0.1 to 5% by weight.

The feed for domestic fowls can be obtained by adding the indigestible dextrin to a basal feed and mixing them, but the method for the addition thereof is not critical. For instance, the indigestible dextrin may be added by distributing or spraying a liquid containing the indigestible dextrin on the basal feed; or mixing the basal feed with the added indigestible dextrin in a powdery state, with the latter method being preferred because it is easily practicable. The feed for domestic fowls of the present invention can be used irrespective of, for in-

stance, the kinds of domestic fowls, daily ages, methods for feeding and breeding and accordingly can be applied to the usual poultry farming method.

The feed for domestic fowls of the present invention is effective over the entire breeding period of the fowl, can prevent, in particular, accumulation of fats in the fowl body and hence formation of fatty liver to thus improve the meat quality, to reduce the content of cholesterol in eggs and to improve the egg-laying rate and the egg-shell strength.

The present invention will hereunder be explained in more detail with reference to the following non-limitative working Examples and the effects practically attained by the invention will also be discussed in detail.

In the following Examples, the content of the indigestible components in the indigestible dextrin, the cholesterol contents in the liver and fowl eggs (yolk) and the triglyceride content in the liver are determined by the methods detailed below, respectively.

[Determination of Content of Indigestible Components in Indigestible Dextrin]

The content of the indigestible components present in the indigestible dextrin used in the feed for domestic fowls is determined by a modified method of that disclosed in "Quantitative Analysis of Indigestible Components" (DENPUN KAGAKU (Starch Science), 1990, Vol 37, No. 2, p. 107).

The modified method comprises accurately weighing out 1 g of a sample of the indigestible dextrin and adding 50 ml of a 0.05 M phosphate buffer (pH 6.0) and then 0.1 ml of α -amylase (Termamy® 120L, having a titer of 120 KNU/g; available from Novo Nordisk Bioindustries, Ltd.) to the sample to react them at 95°C for 30 minutes. After cooling the reaction system, the pH thereof is again adjusted to 4.5, then 0.1 ml of amyloglucosidase (No. A-3042, having a titer of 6100 units/ml; available from Sigma Company) is added to the reaction system, the reaction is continued at 60 °C for 30 minutes and the system is heated up to 90 °C to terminate the reaction. After completion of the reaction, the reaction solution is filled up to 100 ml with water and subjected to determination of the content of glucose present therein by the pyranose oxidase method and thus the content of the indigestible components (%) is calculated from the resulting glucose content (B) and that of the sample (A), separately determined in the same manner, prior to the reaction according to the following equation:

$$\text{The content of indigestible components (\% by weight)} = [1 - A - (B - A) \times 0.9] \times 100$$

wherein A is the glucose content (g) determined prior to the reaction and B is the glucose content (g) determined after the reaction.

[Quantitative Analysis of Cholesterol in Liver]

This method comprises accurately weighing out about 0.1 g of tissue slices of a liver, introducing them into a test tube, adding 2 ml of a 45% by weight aqueous potassium hydroxide solution, heating at 120 °C for one hour in an autoclave, then cooling, adding 2 ml of ethanol, sufficiently stirring in a homomixer, adding 5 ml of n-hexane, stirring for additional 2 minutes, dispensing 1 ml of the hexane phase into another test tube, evaporating the hexane, adding 0.5 ml of isopropyl alcohol, quantitatively determining the amount of cholesterol by the enzyme method and calculating the cholesterol content in the liver sample according to the following equation:

$$\text{Cholesterol Content in Liver (mg/g)} = [\text{measured value (mg/dl)} \times 0.5 \times 5] / [100 \times \text{weight of the liver sample (g)}]$$

[Quantitative Analysis of Triglyceride in Liver]

This method comprises accurately weighing out about 0.1 g of liver tissue slices which are cut into tiny pieces with a razor blade, introducing them into a test tube, adding 5 ml of isopropyl alcohol, allowing to stand for 5 minutes, stirring for 3 minutes in a homomixer to extract lipids present therein, centrifuging 10 minutes at 3000 rpm, quantitatively determining the amount of triglyceride present in the supernatant liquid by the enzyme method and calculating the triglyceride content in the liver sample according to the following equation:

$$\text{Triglyceride Content in Liver (mg/g)} = [\text{measured value (mg/dl)} \times 5] / [100 \times \text{weight of the liver sample (g)}]$$

[Quantitative Analysis of Cholesterol in Fowl Egg]

The yolk is weighed, lyophilized and then pulverized. At this stage, the loss in weight on drying is recorded. The lyophilized sample (about 0.5 g) and a sand (1 g) are introduced into a 50 ml volume graduated flask. To the mixture, there are added 20 ml of a 0.5 mole/l alcoholic KOH solution and 10 ml of isopropyl alcohol,

followed by heating at 60 °C for 80 minutes in warmed water, cooling down to room temperature, addition of isopropyl alcohol to a total volume of 50 ml, filtration, quantitative inspection of the resulting transparent solution as a specimen for the amount of cholesterol present therein and calculating the cholesterol content in the sample according to the following equation:

$$\text{Cholesterol Content in Yolk (mg/g)} = [\text{measured value (mg/dl)}] \times 49.6/E \times (100 - M)/100$$

wherein E means the weight of the lyophilized sample (g) and M means the loss in weight on drying (%).

Example 1

In this Example, fowls belonging to 3-way cross fowls [white Plymouth rock species ♂ × (Satsuma fowl ♂ × Nagoya Species ♀)] which were 110-day-old and which had been put to grassland were used as sample fowls. Ten female fowls were divided into two groups each comprising 5 fowls and each fowl was fattened in an open cage-single pen for 14 days. These fowls of each group were fattened up by feeding, during the fattening period, them with a feed of the present invention comprising a feed for broiler finishing (trade name: Super Final; available from Showa Sangyo Co., Ltd.) having a mixing ratio shown in Table 1 and a composition shown in Table 2 and 5% by weight of an indigestible dextrin B (Fibersol #2; having an indigestible component-content of about 90% by weight) or the feed for broiler finishing per se (control group) for the purpose of comparison.

Table 1

Kinds of Raw Materials	Mixing Ratio	Raw Materials
Cereals	70%	corn, milo
Vegetable Oil Meal	17%	soybean oil meal, corn gluten meal
Animal Feed	8%	fish meal, meat/bone meal
Other Materials	5%	animal oil and fats, lecithin for feed, common salt, calcium

carbonate

5 Feed Additives vitamin A, vitamin D₃, vitamin E,
 10 pantothenic acid, folic acid,
 15 vitamin B₁₂, vitamin K₃, vitamin
 20 B₁, vitamin B₂, vitamin B₆, nico-
 tinic acid, choline, manganese
 sulfate, iron sulfate, copper
 sulfate, zinc sulfate, cobalt
 sulfate, potassium iodate, methi-
 onine, lysine, ethoxyquin

Table 2

Component	Amount
Crude Protein	not less than 18.0%
Crude Fat	not less than 5.0%
Crude Fiber	not more than 4.0%
Crude Ash	not more than 7.0%
Calcium	not less than 0.70%
Phosphorus	not less than 0.55%
Metabolic Energy/Kg	not less than 3200 Kcal

40 After completion of the fattening period, the fowl samples were inspected for the gain of body weight, the amount of feed intake, the weight of the liver, the rate (%) of the liver weight with respect to the body weight, the degree of fatty liver (Fat Liver Score: FLS; 5-stage evaluation according to the color model) and the triglyceride value (TG) in the serum. The results thus obtained are summarized in the following Table 3.

Table 3

Items Determined	Feed of the Invention	Comparative Feed
Gain of Body Weight (g)	432	434
Amount of Feed Intake (g)	2000	2000
Weight of Liver (g)	60.6	72.4
Rate of Liver Weight/Body Weight (%)	2.2	2.5
FLS	3.6	4.2
TG in Serum (mg/dl)	476	671

As seen from the results listed in Table 3, the group fed with the indigestible dextrin-containing feed of the invention had a low liver weight, hence a low rate (%) of the liver weight with respect to the body weight, a low FLS value and hence a low degree of fat-accumulation and a low triglyceride value in the serum as compared with those observed for the group fed with the usual feed free of the indigestible dextrin although the gain of body weight and the amount of feed intake were identical to those for the control group. The low triglyceride value in the serum indicates that the lipid-metabolism of the fowls was improved and that the indigestible dextrin permitted the prevention of the fowls from suffering from hyperlipidemia.

Example 2

In this Example, fowls belonging to 3-way cross fowls [white Plymouth rock species ♂ × (Satsuma fowl ♂ × Nagoya Species ♀)] which were 28-day-old and which had been crate-fattened were used as sample fowls. Male or female chicks (20 chicks each) were divided into two groups each comprising 10 chicks and fattened according to the windowless-floor feeding for 84 days. These chicks of each group were fattened up by feeding, during the fattening period, them with a feed of the present invention comprising a feed for the latter period-broiler (trade name: Super Pro A; available from Showa Sangyo Co., Ltd.) having a mixing ratio shown in Table 4 and a composition shown in Table 2 or a feed for broiler finishing used in Example 1 and 5% by weight of an indigestible dextrin B (Fibersol #2) or either of the feeds free of the indigestible dextrin (control groups) for the purpose of comparison.

Table 4

25	Kinds of Raw	Mixing	Raw Materials
	<u>Materials</u>	<u>Ratio</u>	
	Cereals	70%	corn, milo
30	Vegetable Oil Meal	17%	soybean oil meal, corn gluten meal
	Animal Feed	8%	fish meal, meat/bone meal
35	Other Materials	5%	animal oil and fats, lecithin for

5 Feed Additives feed, common salt, calcium
carbonate
10 nosiheptide, sodium salinomycin,
vitamin A, vitamin D₃, vitamin E,
pantothenic acid, folic acid,
15 vitamin B₁₂, vitamin K₃, vitamin
B₁, vitamin B₂, vitamin B₆, nico-
tinic acid, choline, manganese
carbonate, iron sulfate, copper
20 sulfate, zinc carbonate, cobalt
sulfate, calcium iodate, methio-
nine, lysine, ethoxyquin

25 After completion of the fattening period, the fowl samples were inspected for the gain of body weight, the
amount of feed intake, the weight of the liver, the rate (%) of the liver weight with respect to the body weight,
the degree of fatty liver (Fat Liver Score: FLS; 5-stage evaluation according to the color model, b value deter-
30 mined by a color difference meter), the triglyceride value (TG) in the serum and the fat contents in the liver,
breast meat and dark meat (Soxhlet: SL value; EM scan: EM value). The results thus obtained are summarized
in the following Table 5.

35 Table 5

40 Items Determined Feed of the Comparative

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	<u>Invention</u>	<u>Feed</u>
	Gain of Body Weight (g)	2878
5	Amount of Feed Intake (g)	8634
	Weight of Liver (g)	52.0
10	Rate of Liver Weight/Body Weight (%)	1.7
	FLS	2.4
	b Value	9.8
15	TG in Serum (mg/dl)	31
	Fats in Liver (%)	4.7
	Soxhlet Value (%)	
20	breast meat with the skin	13.0
	dark meat with the skin	18.5
25	EM Scan Value (%) *1	
	breast meat with the skin	14.3
	dark meat with the skin	18.0
30	*1: Calculated on the basis of a regression formula.	

As seen from the results listed in Table 5, the group fed with the indigestible dextrin-containing feed of the invention had a low liver weight, hence a low rate (%) of the liver weight with respect to the body weight, low FLS and b values and hence a low degree of fatty liver and a low triglyceride value in the serum as compared with those observed for the control groups fed with the usual feed free of the indigestible dextrin although the gain of body weight and the amount of feed intake are identical to those for the control group.

Male and female fowls fattened in Example 2 (5 animals each) were inspected for the amounts of cholesterol and triglyceride present in the liver. The results thus obtained are summarized in the following Table 6.

Table 6

Items Determined		Feed of the Invention	Comparative Feed	
Male Fowls				
Cholesterol	mg/g	3.12*	3.54	
	mg/liver	156*	208	
	Triglyceride	mg/g	15.9	18.7
		g/liver	0.91	1.10
Female Fowls				
Cholesterol	mg/g	2.97	2.63	
	mg/liver	141	128	
	Triglyceride	mg/g	31.2*	100.4
		g/liver	1.47*	5.74

The results listed in Table 6 indicate that the male group fed with the feed of the present invention and fattened in Example 2 exhibited a significant reduction of the cholesterol content in the liver, but did not show any significant change in the amount of triglyceride, while the female fowl group fed with the feed of the present invention did not show any significant change in the cholesterol content in the liver, but showed a significant reduction in the triglyceride content.

Then the quality of chicken was evaluated by sensory tests. Dark meat of two female fowls of each group was cut into about 2 cm square cubes, common salt was added in an amount of 1% per unit weight of the meat, heated on a frying pan till even the meat core changed its color to give a meat sample broiled with salt. Separately, breast meat of two male fowls selected from each group was cut into about 2 cm square cubes, immersed in a 5% aqueous common salt solution for 5 minutes and then steamed for 30 minutes to give a salted and steamed sample. The quality of the meat sample broiled with salt was examined by a sensory test whose panel comprised 23 panelists and evaluated according to 5-stage evaluation criteria and the results obtained were summarized in the following Table 7. On the other hand, the salted and steamed sample was likewise examined by a sensory test whose panel comprised 17 panelists and evaluated according to 5-stage evaluation criteria and the results obtained were summarized in the following Table 8.

Table 7

Items Examined	Feed of the	Comp.
	Invention	Feed
Good smell	0%	0%
Palatability	26.9%	17.4%
Smooth and pleasant on the palate	11.5%	0%
Plain taste	3.8%	17.4%

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	Firm and resistant to the teeth	23.1%	8.7%
	Being in prime of grease	3.8%	13.0%
5	Giving out a bad smell	3.8%	17.4%
	Hard	3.8%	4.3%
	Lacking in moisture	19.2%	8.7%
10	Soft	0%	8.7%
	Others	4.1%	4.4%
15	Total Score	76	68.5
	Average	3.6 ±1.2	3.3±1.2

Table 8

25	Items Examined	Feed of the Invention	Comp. Feed
30	Good smell	0%	0%
	Palatability	0%	0%
	Smooth and pleasant on the palate	0%	0%
35	Plain taste	13.3%	18.8%
	Firm and resistant to the teeth	6.7%	25.0%
40	Being in prime of grease	0%	0%
	Giving out a bad smell	0%	12.5%
	Hard	6.7%	18.8%
45	Lacking in moisture	40.0%	12.5%
	Soft	26.7%	6.3%
50	Others	6.6%	6.1%
	Total Score	51.5	51.0
55	Average	3.0 ±0.8	3.0±1.12

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Chicks of ovum recovery-fowls (belonging to DEKALB XL-L(Delta); 24 chicks) were divided into two groups. The first group was fed with a commercially available formula feed (formula feed as a chick starter; available from Nippon Formula Feed Mfg. Co., Ltd.) having a mixing ratio shown in Table 9 and a composition shown in Table 10 up to 4-week-old; with a commercially available formula feed (formula feed for growing middle chicks; available from Nippon Formula Feed Mfg. Co., Ltd.) having a mixing ratio shown in Table 11 and a composition shown in Table 12 during 5 to 10-week-old; with a commercially available formula feed (formula feed for feeding large chicks; available from Nippon Formula Feed Mfg. Co., Ltd.) having a mixing ratio shown in Table 13 and a composition shown in Table 14 during 11 to 20-week-old; and with a commercially available formula feed (formula feed for mature fowls; available from Nippon Formula Feed Mfg. Co., Ltd.) during 21 to 27-week-old.

On the other hand, the second group was fattened up to 27-week-old by feeding them with the foregoing feeds each of which comprised 5% by weight of the indigestible dextrin B (Fibersol #2).

Table 9

5	Kinds of Raw	Mixing	Raw Materials
	<u>Materials</u>	<u>Ratio</u>	
	Cereals	54%	corn, milo
10	Vegetable Oil Meal	26%	soybean oil meal, corn germ meal, rapeseed oil meal
15	Animal Feed	7%	fish meal, meat/bone meal
	Chaff and Bran	7%	corn gluten feed
	Other Materials	5%	animal oil and fats, alfalfa meal, 20 corn steep liquor, calcium carbo- nate, common salt
25	Feed Additives		amprolium, ethopavate, virginia- mycin, vitamin A, vitamin D ₃ , vitamin E, vitamin K ₃ , vitamin B ₁ , 30 vitamin B ₂ , vitamin B ₆ , nicotinic acid, pantothenic acid, biotin, 35 folic acid, vitamin B ₁₂ , choline, methionine, zinc carbonate, calci- um iodate, cobalt sulfate, iron 40 sulfate, copper sulfate, manganese sulfate, ethoxyquin

Table 10

50	<u>Component</u>	<u>Amount</u>
	Crude Protein	not less than 20.3%
	Crude Fat	not less than 2.5%
55	Crude Fiber	not more than 6.0%

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Crude Ash	not more than 8.0%
Calcium	not less than 0.70%
Phosphorus	not less than 0.55%
Metabolic Energy/Kg	not less than 2900 Kcal

Table 11

Kinds of Raw Materials	Mixing Ratio	Raw Materials
Cereals	60%	corn, milo
Vegetable Oil Meal	16%	soybean oil meal, corn germ meal, rapeseed oil meal
Animal Feed	5%	fish meal, meat/bone meal
Chaff and Bran	14%	corn gluten feed, rice bran
Other Materials	5%	animal oil and fats, alfalfa meal, corn steep liquor, calcium carbonate, common salt
Feed Additives		amprolium, ethopavate, sulfaquinoxaline, virginiamycin, vitamin A, vitamin D ₃ , vitamin E, vitamin K ₃ , vitamin B ₁ , vitamin B ₂ , vitamin B ₆ , nicotinic acid, pantothenic acid, biotin, folic acid, vitamin B ₁₂ , choline, methionine, zinc carbonate, calcium iodate, cobalt sulfate, iron sulfate, copper sulfate, manganese sulfate, ethoxyquin

Table 12

Component	Amount
Crude Protein	not less than 17.0%
Crude Fat	not less than 2.5%
Crude Fiber	not more than 6.0%
Crude Ash	not more than 9.0%
Calcium	not less than 0.70%
Phosphorus	not less than 0.55%
Metabolic Energy/Kg	not less than 2770 Kcal

Table 13

5	Kinds of Raw	Mixing	Raw Materials
	<u>Materials</u>	<u>Ratio</u>	
	Cereals	62%	corn, milo, cassava meal
10	Chaff and Bran	15%	corn gluten feed, rice bran, wheat bran
15	Vegetable Oil Meal	11%	soybean oil meal, corn germ meal, rapeseed oil meal
	Animal Feed	5%	fish meal, meat/bone meal
20	Other Materials	7%	alfalfa meal, molasses, corn steep liquor, calcium carbonate, common salt
25	Feed Additives		vitamin A, vitamin D ₃ , vitamin E, vitamin K ₃ , vitamin B ₁ , vitamin B ₂ , vitamin B ₆ , nicotinic acid, panto- thenic acid, biotin, folic acid, vitamin B ₁₂ , choline, methionine, zinc carbonate, calcium iodate, cobalt sulfate, iron sulfate, copper sulfate, manganese sulfate, ethoxyquin
30			
35			
40			
45			
50			
55			

Table 14

Component	Amount
Crude protein	not less than 14.0%
Crude Fat	not less than 2.5%
Crude Fiber	not more than 6.0%
Crude Ash	not more than 9.0%
Calcium	not less than 0.70%
Phosphorus	not less than 0.45%
Metabolic Energy/Kg	not less than 2730 Kcal

After starting the fattening, egg laying was initiated at about 20 weeks. Thereafter, the fowls were inspected for the rate of egg laying (number of eggs produced per day/number of bred fowls) up to 27 weeks at which the rate of egg laying was maximized, the average weight of the eggs produced, the rate of the required feed (total consumption of a feed for mature fowls/total weight of eggs produced), the eggshell strength (the magnitude of the load required for breaking each eggshell) and the content of cholesterol (mg/100 g yolk). The results thus obtained are summarized in the following Table 15.

Table 15

Items Determined	Feed of the Invention	Comparative Feed
Rate of Egg Laying (%)	92.9	84.5
Average Weight of Egg (g)	54.6	53.8
Rate of Required Feed (%)	2.90	2.97
Eggshell Strenth (Kg/cm ²)	3.89	3.40
Content of Cholesterol (mg/100g yolk)	1394	1483

As seen from the results listed in Table 15, the group fed with the indigestible dextrin-containing feed of the invention had a rate of egg laying which was about 8% higher than that observed for the group fed with the comparative feed, a significantly increased eggshell strength ($P < 0.05$) and a significantly reduced cholesterol content ($P < 0.05$).

Example 4

Fowls for ovum recovery (belonging to DEKALB XL-L (Delta); 36 fowls) of 33-week-old were divided into three groups. These fowls were bred for 6 weeks by feeding the first group with a commercially available formula feed for mature fowls (available from Nippon Formula Feed Mfg. Co., Ltd.), feeding the second group with the same feed to which the indigestible dextrin B (Fibersol #2) was added in an amount of 2.5% by weight and feeding the third group with the same feed to which the indigestible dextrin B (Fibersol #2) was added in an amount of 5% by weight. The cholesterol contents were determined before the breeding and 6 weeks after the breeding. The results thus obtained are summarized in Table 16.

Table 16:

Cholesterol Contents (mg/100g yolk)		
Group	Before Breeding	After 6 Weeks
Group 1 (control group)	1435	1556
Group 2 (2.5% dextrin)	1435	1411
Group 3 (5.0% dextrin)	1428	1440

The results listed in Table 16 indicate that the groups fed with the indigestible dextrin-containing feeds had significantly reduced cholesterol contents ($P < 0.05$) in the eggs as compared with the group fed with the usual feed.

Claims

1. A feed for domestic fowls comprising an indigestible dextrin.
2. The feed for domestic fowls of claim 1 wherein the indigestible dextrin is prepared by roasting a starch in the presence of a mineral acid to give a pyrodextrin and treating the pyrodextrin with α -amylase.
3. The feed for domestic fowls of claim 2 wherein the content of the indigestible dextrin ranges from 0.2 to 10% by weight.
4. The feed for domestic fowls of claim 2 wherein the content of indigestible components in the indigestible dextrin ranges from 40 to 60% by weight.
5. The feed for domestic fowls of claim 2 wherein the mineral acid is hydrochloric acid.
6. The feed for domestic fowls of claim 1 wherein the indigestible dextrin is prepared by roasting a starch in the presence of a mineral acid to give a pyrodextrin, treating the pyrodextrin with α -amylase, and then with glucoamylase, filtering and purifying the product, and then separating and removing a digestible fraction from the product through chromatography with an ion exchange resin.
7. The feed for domestic fowls of claim 6 wherein the content of indigestible components in the indigestible dextrin ranges from 50 to 90% by weight.
8. The feed for domestic fowls of claim 6 wherein the content of the indigestible dextrin ranges from 0.1 to 5% by weight.
9. The feed for domestic fowls of claim 6 wherein the pyrodextrin is treated with glucoamylase containing a small amount of α -amylase.
10. The feed for domestic fowls of claim 6 wherein the ion exchange resin is a strongly acidic cation exchange resin which is converted into an alkali or alkaline earth metal form prior to use.



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(54) **Feed for domestic fowls.**

(57) A feed for domestic fowls which comprises an indigestible dextrin prepared by roasting a starch in the presence of a mineral acid to give a pyrodextrin and treating the pyrodextrin with α -amylase or an indigestible dextrin prepared by further treating the α -amylase-treated product with glucoamylase, followed by filtration, purification and separation and removal of digestible fractions of the product through chromatography using an ion exchange resin. The feed serves to inhibit the formation of fatty liver, to improve the meat quality and the rate of egg laying, to improve the eggshell strength and to reduce the cholesterol content in the egg.

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EUROPEAN SEARCH REPORT

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DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. CL.5)
X	EP-A-0 470 895 (MATSUTANI CHEMICAL INDUSTRIES CO. LTD.) * page 3, line 49 - line 50 * * page 4, line 20 - line 49 * * claims 1-4 *	1-10	A23K1/16 A23K1/18
P,X	EP-A-0 535 627 (MATSUTANI CHEMICAL INDUSTRIES CO. LTD.) * examples 1-4, 57 * * claims 1-8, 13, 14 *	1-10	
P,X	EP-A-0 540 421 (MATSUTANI CHEMICAL INDUSTRIES CO. LTD.) * examples 1-4, 57 * * claims 1-7, 13, 14 *	1-10	
X	EP-A-0 530 111 (MATSUTANI CHEMICAL INDUSTRIES CO. LTD.) * examples 6, 60 *	1	
P,X	EP-A-0 538 146 (MATSUTANI CHEMICAL INDUSTRIES CO. LTD.) * examples 6, 60 *	1	TECHNICAL FIELDS SEARCHED (Int. CL.5) A23K
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 16 December 1994	Examiner Dekeirel, M
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